

L 10883-65

ACCESSION NR: AR4046547

the foreign ions cease to participate in the conductivity. The increase in  $\sigma$  with increasing integral radiation dose can be attributed to the formation of additional vacancies (RZhFiz, 1955, no. 10, 23112).

SUB CODE: SS

ENCL: 00

Card 2/2

ACCESSION NR: AP4036568

S/0139/64/000/002/0131/0135

AUTHOR: Pershits, Ya. N.

TITLE: On high temperature electrical conductivity in alkali-halide crystals with impurities

SOURCE: IVUZ. Fizika, no. 2, 1964, 131-135

TOPIC TAGS: electric conductivity, alkali halide, vacant point, lattice, absolute zero, potential barrier, activation energy, ion capture, potassium chloride, potassium bromide

ABSTRACT: Electric conductivity expressed as a function of temperature in alkali-halide crystals KCl and KBr with electrolytically added  $\text{Cu}^+$  ions was determined. Analytically, this is expressed by

$$\sigma = Ae^{-\frac{\epsilon R + \epsilon}{kT}} - Be^{-\frac{\epsilon + \epsilon_1}{kT}}$$

where  $\epsilon$  - work necessary to create a pair of vacant points in a lattice at absolute zero temperature,  $\epsilon_1$  - height of potential barrier for lattice ions at

Card 1/2

ACCESSION NR: AP4036568

absolute zero. Experimental graphs were plotted for  $\sigma = f(1/T)$ , and the activation energy  $E$  of both KCl and KBr was determined from these graphs. In electron-volt units,  $E$  is 2.06 for KCl and  $E$  is 1.97 for KBr ( $E = E/2 + \omega_1$ ). It is shown that the increase in electric conductivity of the alkali-halide crystals is caused by  $\text{Cu}^+$  ion capture in free lattice points. Orig. art. has: 6 formulas, 4 figures, and 1 table.

ASSOCIATION: Pskovskiy gospedinstitut imeni S. M. Kirova (Pskov State Teachers College)

SUBMITTED: 17Mar62

DATE ACQ: 05Jun64

ENCL: 00

SUB CODE: SS

NO REF SOV: 001

OTHER: 006

Card 2/2

ACCESSION NO: A04034661

S/0196/64/000/003/EO05/EO05

SOURCE: Ref. Zh. Elektrot. khim. i energ., Abs. 3320

AUTHOR: Pershita, Ya. N.

TITLE: Effect of foreign ions on the electric conductivity of alkali-halogen crystals. Abstract.

CYBER SOURCE: Izv. Leningr. elektrotekhn. in-ta, vy\*p. 51, 1963, 186

TOPIC TAGS: ionic crystal, crystal conductivity, foreign ions in crystal, alkali halogen crystal

TRANSLATION: On electrolytic introduction of Cu-ions into alkali-halogen crystals (KCl, KBr), a parallelism can be seen between the logarithmic lines of temperature vs. conductivity for pure crystals and those for the crystals with impurity introduced up to the saturation point at the measurement temperature. Thus, for the high-temperature impurity conductivity, the activation energy coincides with that of majority carriers. The position of the front of the ion cloud leaving the crystal and the current measured in the crystal permit determining the mobility and concentration of foreign ions. The activation-energy components are determined from the temperature-conductivity curve and the mobility. [Pskovskiy gos. pedagogich. in-t im. S. M. Kirova]

1/1 DATE ACQ: 10Apr64

SUB CODE: # SS

ENCL: 00

OCT 19 1964

Aerospace Information



PERSHITS, Ya.N.; GUTMAN, V.I.; ANOSOVA, A.I.

Effect of foreign ions on the electroconductivity of irradiated  
alkali halide crystals. Izv. vys. ucheb. zav.; fiz. no.1:3-7  
'64. (MIRA 17:3)

1. Pskovskiy gosudarstvennyy pedagogicheskiy institut imeni  
Kirova.

PERSHITS, Ya.N.

Motion of metal ions in alkali halide crystals. Uch.zap.Ped.inst.Gerts.  
no.207:132-148 '61.

(MIRA 16:5)

1. Pskovskiy gosudarstvennyy pedagogicheskiy institut imeni S.M.  
Kirova.

(Alkali metal halide crystals)

(Metal ions)

PERSHITS, Ya.N.; GUTMAN, V.I.

Effect of capture centers on the electric properties of alkali halide crystals. Uch.zap.Ped.inst.Gerts.no.207:149-162 '61.

(MIRA 16 5)

1. Pskovskiy gosudarstvennyy pedagogicheskiy institut imeni S.M. Kirova.

(Electrons—Capture)

(Alkali metal halide crystals—Electric properties)



PEESHITS, Ya. N.

Effect of strange ions on the electroconductivity of alkali  
metal halide crystals. Izv. vys. uch. zav., fiz. 3:3-8 '62.  
(MIRA 15:10)

1. Pskovskiy pedagogicheskiy institut imeni S. M. Kirova.

(Alkali metal halide crystals—Electric properties)  
(Ions)

L 13034-63 EPF(n)-2/EWP(q)/EWT(m)/BDS/T-2 AFPTC/ASD/SSD Pu-4

JD/WH

ACCESSION NR: AP3000613

S/0181/63/005/005/1348/1357

AUTHOR: Pershits, Ya. N.

TITLE: The mechanism of changing conductivity during aging of rutile ceramics and the electrolysis of alkali-halide crystals

SOURCE: Fizika tverdogo tela, v. 5, no. 5, 1963, 1348-1357

TOPIC TAGS: rutile, titanium, tantalum, alkali halide, ceramic aging, regeneration, F-center, Cu, KBr, dislocation, conductivity, rutile ceramic, rutile ceramic aging

ABSTRACT: The author's purpose is to show experimentally that not only regeneration but also succeeding aging of Ti-bearing ceramics conforms to the same pattern that is observed in alkali-halide crystals during comparable methods of investigation. In his experiment he used the staining method with a flat tantalum electrode, and found great similarity between the Ti-bearing ceramics and the alkali-halides. His results show that at relatively low temperatures (100-300°C) the movement of added current carriers takes place within the crystal grains of rutile or within mosaic blocks, forming dislocations that remain in the crystal after annealing. These carriers could be F-centers, but since such

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L 13034-63

ACCESSION NR: AP3000613

centers may be displaced only when ionic conductivity is present, aging clearly begins at temperatures giving rise to marked displacement of ions in the electrical field. The identical character of the time relationship of current at all temperatures during which aging was observed clearly points to electrolytic processes in association with change from regeneration to renewed aging even at low temperatures. Considerable doubt is cast on data suggesting the electrolytic conductivity is absent in titanium oxide. It is more likely that the electrolytic products gather at the boundaries of mosaic blocks or crystal grains. The aging of Ti-bearing ceramics at high temperatures is associated with the spread of current carriers from both electrodes, but regeneration is associated with the movement of electron clouds. Orig. art. has: 8 figures, 2 tables, and 11 formulas.

ASSOCIATI: Pskovskiy gosudarstvennyy pedagogicheskiy institut im. S. M. Kirova  
(Pskov State Pedagogical Institute)

SUBMITTED: 09Nov62

DATE ACQ: 11Jun63

ENCL: 00

SUB CODE: 00

NO REF SOV: 012

OTHER: 004

Card 2/2

PERSHITS, Ya.N.

Mechanism underlying conductivity variations in the aging of  
rutile ceramics and the electrolysis of alkali halide crystals.  
Fiz.tver.tela 5 no.5:1348-1357 My '63. (MIRA 16:6)

1. Pakovskiy gosudarstvennyy pedagogicheskiy institut imeni  
Kirova.

(Rutile--Electric properties) (Alkali metal halide crystals)

PERSHITS, Ya.N.; GUTMAN, V.I.

Effect of foreign ions on electron processes in ionic crystals.  
Izv.vys.uch.zav.; fiz. no.4:123-129 '62. (MIRA 15:9)

1. Pskovskiy gosudarstvennyy pedagogicheskiy institut.  
(Ionic crystals) (Electrons)

S/139/62/000/004/010/018  
E132/E435

AUTHORS: Pershits, Ya.N., Gutman, V.I.

TITLE: The influence of foreign ions on electronic processes  
in ionic crystals

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy. Fizika,  
no.4, 1962, 123-129

TEXT: It is established that the influence of added colouring on the electrical properties of alkali halide crystals is caused by the presence in them as impurities of heavy metals, the ions of which act as centres for trapping electrons. The trapping of electrons by foreign ions in crystals irradiated by X-rays has been studied experimentally. A means of colouring crystals from a plane metallic cathode is described. Trapping of electrons by ionic impurities leads to additive or subtractive colouring of the crystal. As a result of additive colouring the neutralization of impurity ions takes place decreasing the number of trapping centres. The trapping of electrons leads to changes in the electrical properties of the crystals. There are 6 figures.

ASSOCIATION: Pskovskiy gospedinstitut (Pskov State-Pedagogical  
Institute)

SUBMITTED: May 3, 1961  
Card 1/1

24.7700

39031  
S/139/62/000/003/001/021  
E039/E420

AUTHOR: Pershits, Ya.N.

TITLE: The influence of impurity ions on the electrical conductivity of alkali-halide crystals

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy. Fizika, no. 3, 1962, 3-8

TEXT: Copper ions are introduced into KBr crystals by passing an electric current through the crystal using a copper anode. The temperature dependence of the electrical conductivity of KBr is measured for different temperatures of introduction of copper and the activation energy is shown to be  $\sim 0.1$  ev less than for the pure crystal. There is a significant increase in conductivity as the  $\text{Cu}^+$  ions enter the crystals and a corresponding decrease when they are removed by reversing the direction of the electric field. Measurements are made of the mobility of copper ions in KBr, KCl and NaCl. The temperature dependence of the concentration of copper ions coincides with the temperature dependence of ionic vacancies in the crystals (at  $600^\circ\text{C}$  the concentration of ions is  $\sim 5 \times 10^{17}\text{cm}^{-3}$ ).  
Card 1/2

PERSHITS, Ya.N.

Aging and regeneration of ceramics containing titanium. Fiz.tver.  
tela 3 no.11: 495-3502 N '61. (MIRA 14:10)

1. Pskovskiy gosudarstvennyy pedagogicheskiy institut im.  
M.Kirova.

(Ceramics) (Titanium)



15.2640

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161/003/011/036/056

7  
C 13138

AUTHOR: Pershits, Ya. N.

TITLE: Ageing and recovery of titanium containing ceramics

PERIODICAL: Fizika tverdogo tela, v. 3, no. 1, 1961, 3495-3502

TEXT: The electrical conductivity of ceramic containing titanium increases when a constant electric field is applied for some time. This process is termed ageing. When the direction of the field is reversed, electrical conductivity will rapidly drop again which process is called recovery. The variation of current with time was measured in T-80 (T-80) titkond ceramics in the temperature range 180-1000°C. Disks with graphite electrodes were used as samples. Recovery can be treated theoretically as the migration of an additional-carrier along grain boundaries. This was confirmed by experiments which showed that the mechanism holds good at low temperatures (i.e.  $t < 400^\circ\text{C}$ ). At high temperatures, intrinsic conductivity will become predominant. The logarithm of the current was found to be a linear function of the logarithm of time. However, the original conductivity was reached after recovery. The specimens started ageing again.

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Card 1/2

Ageing and recovery of...

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5/16/61/003/011/036/056  
5104/H138

Comparative measurements with alkali-halide crystals yielded qualitatively the same results as were obtained for titanium-containing ceramics (S. A. Artsybyshch. Tr. Fiz. inst. im. P. N. Lebedeva, 1, 5, 1938). Ageing was found to be due to electron capture by anionic vacancies. Recovery, on the other hand, is due to the formation of F-centers which migrate within the grains. Since the F-centers are partly dissociated into free electrons and lattice defects, recovery is characterized by two processes: one rapid (linear dependence of  $\log I$  on  $\log t$ ) and one slow. There are 5 figures, 1 table, and 10 references: 9 Soviet-bloc and 1 non-Soviet-bloc. The reference to the English-language publication reads as follows: W. A. Weyl and N. A. Terhune. Ceram. Age, 62, 23, 1953.

ASSOCIATION: Pskovskiy gosudarstvennyy pedagogicheskii institut im. S. M. Kirova (Pskovsk State Pedagogical Institute imeni S. M. Kirov)

SUBMITTED: April 15, 1961 (initially)  
June 26, 1961 (after revision)

Card 2/2

PERSHITS, Ya.N.; TARASENKO, G.D.

Effect of additive coloring on electric properties of alkali  
halide crystals. Izv.vys.ucheb.zav.; fiz. no.5:121-126 '58.  
(MIRA 12:1)

1. Pekovskiy pedinstitut i Severo-Osetinskiy pedinstitut.  
(Alkali halide crystals--Electric properties)

AUTHORS: Pershits, Ya. N., Tarasenko, G. D.

TITLE: Influence of Additive Coloration on Electrical Properties of Alkali-Halide Crystals (Vliyaniye additivnogo okrashivaniya na elektricheskiye svoystva shchelocno-galoidnykh kristallov)

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy, fizika, 1958, Nr 5, pp 121-126 (USSR)

ABSTRACT: An alkali-halide crystal lattice is postulated in which the density of 'natural' impurities (F-centres, lattice imperfections) is constant, and into which is introduced an additional type of impurity whose concentration varies in a known manner with time. The added impurity atoms are (or become) ionised and contribute to the electrical properties of the lattice. One may associate for example an ionic conductivity  $\sigma_1$  with the added impurity, which is distinguishable from the ionic conductivity  $\sigma_1$  associated with the F-centres. When a potential difference  $P$  is applied across such a crystal the following relations hold:

$$\frac{dx}{dt} = - \frac{\sigma_1 P}{\sigma_1^2 + \sigma_2 - \sigma_1} \cdot \frac{1}{E^2} \cdot \frac{dE}{dt} = uE \quad (1)$$

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SOV/139-58-5-26/55

# Influence of Additive Coloration on Electrical Properties of Alkali-Halide Crystals

Here  $E$  is the field associated with  $P$ ,  $\sigma_2$  is the electronic conductivity, deriving from the electrons of the added impurity atoms,  $x$  is a measure of the concentration of impurity atoms, and  $u$  is the mobility of the F-centres. The  $\sigma$ 's and  $u$  are related, via the mean free path, to the lattice temperature, and this leads to the following expression for the total current at time  $t$  and temperature  $T$ :

$$I = I_0 I_1 \left[ I_1^2 + (I_0^2 - I_1^2) \frac{t}{T} \right]^{-\frac{1}{2}} \quad (3)$$

where  $I_0$ ,  $I_1$  are respectively the current contributions from the added impurity atoms and the F-centres. If one further denotes the ionic part of  $I_0$  by  $I_{0i}$  (and hence the electronic part by  $I_0 - I_{0i}$ ) then the F-centre concentration may be

Card 2/4

OW/124-50-5-46/55

Influence of Additive Coloration on Electrical Properties of Alkali-Halide Crystals

expressed in terms of the I's thus:

$$N = \frac{2TI_1}{e v} \cdot \frac{I_0 - I_{01}}{I_0 + I_1} \quad (4)$$

where  $e$  denotes the electronic charge and  $v$  the crystal volume. Hence measurements of the impurity current enable the F-centre concentration to be deduced. Further, measurements of  $\sigma$  at various temperatures and known impurity concentrations enable the F-centre mobility to be determined:

$\log \sigma$  plotted against  $T^{-1}$  gives a straight line whose slope is the mobility. Such curves were plotted from measurements on crystals of KCl, KBr and NaCl, to which impurities had been added, at various temperatures. The effect of the added

impurity was to increase the F-centre mobility by a factor varying from about 3 at 600°C to 5 at 450°C. The work was first reported at the Conference of Higher Education Establishments on Dielectrics and Semiconductors at Tomsk, in

Card 3/4

SOV/139-58-5-26/35

Influence of Additive Coloration on Electrical Properties of Alkali-Halide Crystals

February, 1958. The paper contains 1 table, 4 figures and 9 references (5 Soviet, 4 German).

ASSOCIATION: Pskovskiy pedinstitut; Severo-osetinskiy pedinstitut  
(Pskov Pedagogical Institute; North Osetian Pedagogical Institute)

SUBMITTED: April 7, 1958.

Card 4/4

PERSHITS, Ya. N., Docent

Docent Ya. N. Pershits (Pskov Institute of Pedagogics) and G. D. Tarasenko (North Ossetian Institute of Pedagogics, Ordzhonikidze)

"The character of the propagation of the F-center cloudlet and that of its mobility is changed in consequence of the primary coloration of the NaCl-, KCl-, KBr- and KJ crystals, whereas the electric conductivity of the ions of the samples is reduced irreversibly"

Report presented at a Conference on Solid Dielectrics and Semiconductors,  
Tomsk Polytechnical Inst., 3-8 Feb. 58.  
(Elektrichestvo, '58, No. 7, 83-86)



PERSHITS, Ya. N.

Pershits, Ya. N. [Pskovskiy pedagogicheskiy institut (Pskov Pedagogical Institute)] Phenomena in the Conductivity of Dielectrics Formed Near the Cathode

(The Physics of Dielectrics, Conductivity, and the Theory of Dielectrics) Moscow: Nauka, 1984. 114 p. 24 cm. 1.50 rub. 1.50 rub. 1.50 rub.

This volume published in the series "Fundamental Problems of Physics" (Series "Fundamental Problems of Physics") is devoted to the study of the phenomena of dielectric breakdown and the theory of dielectrics. It is a collection of papers presented at the International Symposium on Dielectrics, held in Leningrad in 1982. The papers are devoted to the study of the phenomena of dielectric breakdown and the theory of dielectrics. The papers are presented in the form of a collection of papers presented at the International Symposium on Dielectrics, held in Leningrad in 1982. The papers are devoted to the study of the phenomena of dielectric breakdown and the theory of dielectrics. The papers are presented in the form of a collection of papers presented at the International Symposium on Dielectrics, held in Leningrad in 1982. The papers are devoted to the study of the phenomena of dielectric breakdown and the theory of dielectrics.

*PERSHITS, V. A.*

Distr: 434/4330/4Euc/4E30

Thermal electronic conductivity in alkali halide crystals.  
~~V. A. Pershits, Uchenye Zapiski Pribl. Kazansk. U-~~  
~~chensk. Univ., 1935, No. 3, 210-233; Referat. Zhur., Khim.~~  
~~1936, Abstr. No. 33142.—A review of publications on prop-~~  
~~erties of color centers and dark-cond. of colored crystals.~~  
~~58 references. V. S. Mikhajlov.~~

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SOV/58-59-5-10861

Translation from: Referativnyy Zhurnal Fizika, 1959, Nr 5, p 137 (USSR)

AUTHOR: Pershits, Ya.N.

TITLE: Phenomena Near the Cathode During Electric Conduction in Dielectrics  
That are Being Formed

PERIODICAL: V sb.: Fiz. dielektrikov. Moscow, AS USSR, 1958, pp 76 ~ 82

ABSTRACT: It is shown that when a current is passed through eternit (asbestos cement), the resistance of the contact with the metallic cathode decreases on the average by half in the course of a few seconds and then does not change. If the layer near the cathode is sandpapered, the conductivity of this layer will be found to increase again. A reduction in the electric field leads to relaxation phenomena: if the time-dependence curve of the current is remeasured 24 hours after the first measurement, its shape will be about the same as that of the first measurement, but the conductivity of the contact will have decreased. In the event of prolonged electrolysis in eternit at 18°C, the process of the increase in conductivity of the layer near the cathode is offset by the formation of a layer near the cathode exhibiting unipolar conductivity and high

Card 1/2

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204/55-55-1-55-5

Translation from: Referativnyy Zhurnal Fizika, 1959, Nr 7, pp 145 - 146 (USSR)

AUTHOR: Pershits, Ya.N.

TITLE: Phenomena Pertaining to the Region Near the Electrode in Dielectrics <sup>21</sup>

PERIODICAL: Uch. zap. Leningr. gos. ped. in-ta im. A.I. Gertsena, 1958, Vol 148, pp 131 - 149

ABSTRACT: Using a string electrometer or a cathode oscillograph with photographic registration, the author studied the time dependences of "back" current ( $I_b$ ) and forward current ( $I_f$ ) in quartz, glass, NaCl, KCl, Eternit (asbestos-cement), porcelain, mica and asbestos. In order to eliminate the effect of the second electrode and to make a more detailed study of the formation phenomenon, the author used the auxiliary electrode method (Zh. eksperim. i teor. fiz., 1947, Nr 17, p 251). It was found that with a change in the direction of the field,  $I_b$  falls off according to a curve with an inflection point whose abscissa is proportional to the time of passage of the current in the forward direction. The identical character of the dependence of  $I_b$  in the case of various dielectrics indicates that the generally accepted mechanism of the formation of

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67191

SOV/58-59-7-15678

Phenomena Pertaining to the Region Near the Electrode in Dielectrics

dielectrics (freeing the layer near the electrodes from impurities due to electrolysis) does not explain the observed phenomena. This conclusion is also borne out by a study of the variations that the unipolarity coefficient undergoes when samples are subjected to heat treatment. At low temperatures formation phenomena were observed that resemble those at high temperatures. It is assumed that under the action of the electric field and temperature, phenomena occur in the layers near the electrodes that lead to a change in the resistance and stability of these regions. It is possible that the formation of the anode and cathode layers is due to processes involving anions and cations respectively. The hypothesis that electron conduction takes place throughout the entire volume of a dielectric that is being formed, contradicts a number of experimental data. ✓

L.K.

Card 2/2

GALEEV, A. M., doktor tekhnicheskikh nauk, professor; PARSHITS, Yu. I.

Characteristics of sliding friction during the movement of a body  
along an anisotropic surface. Izv. VNIIT no. 92/11:169-180 '57.

(MLRA 10:7)

(Friction) (Surfaces (Technology))

SOV 124 58 11 12156

Translation from: Referativnyy zhurnal, Mekhanika 1958 Nr 11 p 26 (USSR)

AUTHORS: Galeev, A. U., Pershits, Yu. I.

TITLE: Peculiarities of the Sliding Friction of a Body Against an Anisotropic Surface (Osobennost treniya skorozheniya pri dviizhenii tela po anizotropnoy poverkhnosti)

PERIODICAL: Tr. Mosk. in-ta inzh. zh. d. transp. 1957, Nr 92 11 pp 169-183

ABSTRACT: The authors call attention to the great difference often encountered in the friction coefficients of a surface measured in the direction in which the surface has been machined as against measurements thereof made in other directions. In this connection they pose and proceed to investigate, in a general qualitative form, the problem of the friction of a mass point moving over a surface having anisotropic frictional characteristics. As the basic characteristic of the frictional properties of the surface they introduce a polar curve showing the relationship between the specific work of the force of friction and the direction of travel of the mass point. For the purpose of some computational estimates they assume this curve to be an ellipse, but, in principle, the investigation remains the same whatever the shape of the curve.

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SOV 124 58 11 1956

Peculiarities of the Sliding Friction of a Body (cont.)

Further, they construct on the same scale a curve of the specific work of the force exerted on a point as a function of its direction, this second curve being a circle erected on the vector of the acting force as its diameter. In this event motion may occur in directions in which the second curve is found to extend out beyond the first, and it is the authors' assumption that it will actually occur in that direction in which the difference between the radius-vectors of the two curves is maximal. It is the opinion of this reviewer that the authors' designation of the true direction of motion requires further proof. Therefore, as it turns out, the direction of motion does not, in general, coincide with the direction of the force applied. Assuming the curve of the specific work performed by the force of friction to be an ellipse, the authors are able to determine the angle between the direction of motion and the direction of the acting force. In conclusion, they examine the subject of construction of the cone of friction for an anisotropic surface.

G. I. Barenblatt

Card 2/2



PERSHITS, YU. I. Cand Tech Sci -- (diss) "Investigations of forces occurring in automatic couplers during the braking of loaded trains," Moscow, 1960, 26 pp, 120 cop. (Moscow Institute of Engineers of Railroad Transport im I. V. Stalin) (KL, 42-60, 114)

GALEYEV, A.U., doktor tekhn.nauk, prof.; PERSHITS, Yu.I., inzh.

Determining impact forces occurring in automatic couplings  
during emergency braking of freight trains using 270-002 air  
distributors. Trudy MIIT no.102:5-19 '59. (MIRA 12:10)  
(Car couplings) (Railroads--Brakes)

GALEEV, Akhmet Umerovich, prof.; PERSHITS, Yuliy Isaakovich, inzh.;  
SAZONOV, A.G., inzh., red.; BOBROVA, Ye.N., tekhn. red.

[Problems of the mechanics of trains] Voprosy mekhaniki poezda.  
Moskva, Gos. transp. zhel.-dor. izd-vo, 1958. 231 p. (MIRA 11:9)  
(Railroads--Trains)

GALEEV, Akhmet Umerovich [deceased]; PERSHITS, Yuliy Isaakovich; MELAMED, D.A., inzh., retsenzent; LEBEDEV, A.V., inzh., retsenzent; SOBAKIN, V.V., inzh., red.; BOBROVA, Ye.N., tekhn. red.

[Fundamentals of mechanics for locomotive crews] Osnovy mekhaniki dlia lokomotivnykh brigad. Moskva, Vses.izdatel'sko-poligr.ob"edinenie M-va putei soobshcheniia, 1961. 167 p. (MIRA 14:11)  
(Mechanics) (Railroads)

PHASE I BOOK EXPLOITATION

SOV/5054

Vsesoyuznaya konferentsiya po treniyu i iznosu v mashinakh. 3d, 1958.

Sukhoie i granichnoye treniye. Friksionnyye materialy (Dry and Boundary Friction. Friction Materials) Moscow, Izd-vo AN SSSR, 1960. 302 p. Errata slip inserted. 3,500 copies printed. (Series: Its: Trudy, v. 2)

Sponsoring Agency: Akademiya nauk SSSR. Institut mashinovedeniya. Resp. Ed.: I. V. Kragel'skiy, Doctor of Technical Sciences, Professor; Ed. of Publishing House: K. I. Grigorash; Tech. Ed.: S. G. Tikhomirova.

PURPOSE: This collection of articles is intended for practicing engineers and research scientists.

COVERAGE: The collection published by the Institut mashinovedeniya, AN SSSR (Institute of Science of Machines, Academy of Sciences USSR) contains papers presented at the III Vsesoyuznaya konferentsiya po treniyu i iznosu v mashinakh (Third All-Union

Card 1/11

Dry and Boundary Friction (Cont.)

SOV/5054

Conference on Friction and Wear in Machines, April 9-15, 1958. Problems discussed were in 5 main areas: 1) Hydrodynamic Theory of Lubrication and Friction Bearings (Chairmen: Ye. M. Gut'yar, Doctor of Technical Sciences, and A. K. D'yachkov, Doctor of Technical Sciences); 2) Lubrication and Lubricant Materials (Chairman: G. V. Vinogradov, Doctor of Chemical Sciences); 3) Dry and Boundary Friction (Chairmen: B. V. Deryagin, Corresponding Member of the Academy of Sciences USSR, and I. V. Kragel'skiy, Doctor of Technical Sciences); 4) Wear and Wear Resistance (Chairman: M. M. Krushchov, Doctor of Technical Sciences); and 5) Friction and Antifriction Materials (Chairmen: I. V. Kragel'skiy, Doctor of Technical Sciences, and M. M. Krushchov, Doctor of Technical Sciences). Chairman of the general assembly (on the first and last day of the conference) was Academician A. A. Blagonravov. L. Yu. Pruzhanskiy, Candidate of Technical Sciences, was scientific secretary. The transactions of the conference were published in 3 volumes of which the present is the second. This volume contains articles on friction materials, and on various aspects of dry friction and boundary lubrication. Among the broad areas covered are: the

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Dry and Boundary Friction (Cont.)

SOV/5054

- Galeyev, A. U., and Yu. I. Pershits. Sliding Friction During the Motion of a Material Point Along an Anisotropic Surface 15
- Gal'chenko, V. V., and D. Ya. Speranskiy. Investigation of the Relationship Between the Temperature During Friction and the Physical Properties of the Surface Layers of the Rubbing Components of Machines 22
- Garkunov, D. N. Some Laws of the Wear of Metals Under Conditions of Dry and Boundary Friction and Means for Reducing the Wear 26
- Glagolev, N. I. Work of Friction Forces and Wear of Rotating Bodies 34
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Card 4/11

PERSHEKHA, A.A. (Orshansk)

Acute appendicitis. Fel'd i akush. 23 no.7:12-15 J1'58 (MIRA 11:8)  
(APPENDICITIS)



10

PROCESSED AND PREPARED BY THE

Investigation on corrosion and factors governing the selection of alloys in the construction of equipment for the manufacture of caustic soda and caustic potash. M. V. PARANEK AND (Miss) L. POROVA. *Chemie & Industrie Special No.*, 232-4 (March, 1930). — After a brief review of the literature on corrosion of Fe by alkalis, results of corrosion experiments by NaOH and KOH (850 g. per l. and 12 g. Na<sub>2</sub>CO<sub>3</sub> and NaCl) for 8 hrs at 100–400° are given, showing that, with gray cast Fe, the main factor in resistance to corrosion is homogeneity of structure, and compn. is of secondary importance. Addn. of Ni reduces the corrosion, which is completely stopped with 12% or more of Ni. Addn. of Cr is less effective, particularly against the action of KOH. Addn. of 6% Ni and 5% Cr is equiv. to 12% Ni. On the whole, corrosion by KOH is 2.5–3 times as great as that by NaOH. Cu is satisfactory for the piping and fittings, as far as resistance to corrosion is concerned, but is too soft; the most satisfactory alloy was found to be obtained by addn. of 21% Ni to Cu. Also in *J. Chem. Ind. (Moscow)* 7, 16 20(1930).  
A. PAPINRAU-COUTURE

ASD-SLA METALLURGICAL LITERATURE CLASSIFICATION

PROCESS AND PROPERTY INDEX

4

**Influence of External Factors on the Corrosion of Lead in Sulphuric Acid Production.** V. Pershke and V. Ignatieva (*Khimicheskii Zhurnal*, 1933, 7, 38-40, 6 figs., 1933, 39, 3786).—[In Russian.] (*Chem. Abstr.*, 1934, 1, 419) In the uniformity even and the local disintegrations of lead lining of sulphuric acid chambers and towers, the latter form of corrosion may be ascribed to a formation of a galvanic element: (+)  $PbH_2SO_4$  or nitrous acid |  $H_2SO_4$  (or nitrous acid) -  $HNO_3$ ,  $Pb(-)$ . The mechanical imperfections of lining (folds, seams, &c.) cause local differences in temperature and condensation of the reacting mixture. This contains traces of free nitric acid, as a result of varying conditions of production, which attacks lead. The lead nitrate formed is immediately converted, in the presence of sulphuric acid, into lead sulphate, with liberation of nitric acid, and the cycle is indefinitely repeated. These spongy and brittle corrosion products, deposited on the anode, easily fall off, exposing a new surface to the destructive process. Detection of lead nitrate in the corrosion products substantiates the proposed explanation of the corrosion process. The corrosion can be counteracted by a preliminary exposure of lead lining to the action of sulphur dioxide.—S. G.

METALLURGICAL LITERATURE CLASSIFICATION

ELECTRICITY

| 1ST AND 2ND LETTER                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 3RD LETTER    |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 4TH AND 5TH COULDS |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 6TH GROUP       |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
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| AUTHOR INDEX                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | SUBJECT INDEX |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | CROSS REFERENCE    |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | CROSS REFERENCE |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| <p>Parabke, V., and Lashin, N. ALUMINUM SULFATE: ITS PRODUCTION FROM CLAYS FROM THE DISTRICT OF POLEVSKII / <i>Chem. Ind. (U.S.S.R.)</i>, 7, 654-56 (1930). White clays with an average 34.1% alumina content are treated with sulfuric acid to produce raw aluminum sulfate. The disintegration of uncalcined clay by sulfuric acid of 40 BE is slow and produces only small quantities of aluminum sulfate. On the other hand, with clay calcined at 700° for 2 hr., and then treated with sulfuric acid for 7 hr., the surplus acid being neutralized by a 15% addition of clay, the yield is 87.6% of aluminum sulfate 5 hr. after the beginning of the reaction. The insoluble residue contains 78.6% silica which is used for preparing sodium silicate by boiling it for 1 hr. with caustic soda. The silicate contains about 43% <math>\text{Na}_2\text{SiO}_3</math> or <math>\text{K}_2\text{SiO}_3</math> and an insignificant quantity of free caustic soda.</p> |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |               |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |                    |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |                 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

| TIT AND 2ND EXPD                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |  |  |  |  |  |  |  |  |  | PROCESS AND PROPERTY INDEX |  |  |  |  |  |  |  |  |  | 3RD AND 4TH EXPD |  |  |  |  |  |  |  |  |  |
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| <p>3C</p> <p>Aluminum sulphate from clay of the Polovaki district. Y. F. FOMIN and N. LAMIN (J. Chem. Ind., Moscow, 1950, 7, 884-888).--The clay (34.1% <math>Al_2O_3</math>) is heated at <math>700^\circ</math> for 2 hr. and then treated with <math>H_2SO_4</math> for 7 hr., the excess of acid being neutralized by 15% of clay after 5 hr.; 87.5% of <math>Al_2(SO_4)_3</math> is obtained. The residue contained 78.6% <math>SiO_2</math>.</p> <p>CHEMICAL ABSTRACTS.</p> |  |  |  |  |  |  |  |  |  |                            |  |  |  |  |  |  |  |  |  |                  |  |  |  |  |  |  |  |  |  |
| <p>ABB. 35.4 METALLURGICAL LITERATURE CLASSIFICATION</p>                                                                                                                                                                                                                                                                                                                                                                                                                          |  |  |  |  |  |  |  |  |  |                            |  |  |  |  |  |  |  |  |  |                  |  |  |  |  |  |  |  |  |  |
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~~RESTRICTED~~

PERSHKE, V.

PERSHKE, V.

J. Russ. Phys.-Chem. Soc. 60, 1019-35 (1928);

Z. Elektrochem. 35, 13-7 (1929)

CA: 23-2338/9

The relations between the physical constants of a liquid.

~~RESTRICTED~~

**Influence of external factors on the corrosion of lead in sulfuric acid production.** V. Pershke and V. Ignat'eva.

*Khimicheskii Y. Zh.* (1961), 1, 28, 2831. (Of the uniformly even and the local disintegrations of Pb lining of H<sub>2</sub>SO<sub>4</sub> chambers and towers, the latter form of corrosion may be attributed to a formation of a galvanic element (1) Pb-H<sub>2</sub>SO<sub>4</sub>, or nitrous acid H<sub>2</sub>SO<sub>4</sub>, (or nitrous acid) HNO<sub>3</sub>Pb (-). The mechanical imperfections of lining (folds, seams, etc.) cause local differences in temp. and condensation of the reacting mixt. This contains traces of free HNO<sub>3</sub>, as a result of various conditions of production, which attacks Pb. The Pb(NO<sub>3</sub>)<sub>2</sub> formed is immediately converted, in the presence of H<sub>2</sub>SO<sub>4</sub>, into PbSO<sub>4</sub>, with liberation of HNO<sub>3</sub>, and the cycle is indefinitely repeated. These massive and brittle corrosion products, deposited on the anode, easily fall off, exposing a new surface to the destructive process. Detection of Pb(NO<sub>3</sub>)<sub>2</sub> in the corrosion products substantiates the proposed explanation of the corrosive process. The corrosion can be counteracted by a preliminary exposure of Pb lining to the action of SO<sub>2</sub>. Chas. Blanc

ASH-SLA METALLURGICAL LITERATURE CLASSIFICATION

*CA*

18

Aluminum sulfate obtained from the clay of the Pulovskii district. V. Jansson and N. Larsson. *J. Chem. Ind. (Moscow)* 7, 684-6 (1930). The white clay contg. an av. of 34.1%  $\text{Al}_2\text{O}_3$  was treated with  $\text{H}_2\text{SO}_4$  to produce the standard technical  $\text{Al}(\text{NH}_4)_2$ . Decompos. of non-lignted clay with 40%  $\text{H}_2\text{SO}_4$  progressed very slowly and gave small quantities of  $\text{Al}(\text{NH}_4)_2$ . On the contrary when the clay was lignted at 700° for 2 hrs. and then treated with  $\text{H}_2\text{SO}_4$  for 7 hrs. and the excess of acid was neutralized by the addn. of 15% of clay, at the expiration of 6 hrs. from the start of the reaction, 87.6%  $\text{Al}(\text{NH}_4)_2$  was obtained. The insol. residue contg. 78.6%  $\text{SiO}_2$  could be used for the prepn. of water glass by boiling it with caustic for 1 hr. The silicate so obtained contained about 43%  $\text{Na}_2\text{SiO}_3$  (or  $\text{K}_2\text{SiO}_3$ ) and an insignificant amt. of free caustic. V. D. Karpavich.

ASSOCIATED METALLURGICAL LITERATURE CLASSIFICATION

SEARCHED INDEXED

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PERSHKE, V. K.

✓

Rate of transformation of ammonium nitrate from  $\alpha$ - to  $\beta$ -form. V. K. Pershke and A. M. Conny. J. Gen. Chem. (U. S. S. R.) 7, 303-8 (1937); cf. C. A. 31, 2017 (1937), 2109s. The rate of transformation increases in the presence of moisture and with increase in number of transformation centers. It increases in direct proportion to the difference between the temp. at which transformation is taking place and the true transformation temp. which was found to be 33.0-33.6°. S. L. Alshukh.





MA

*Corrosion and Related Phenomena*

**The Action of Sulphuryl Chloride on Metals.** V. K. Farkas and L. J. Twyman. *Trans. Faraday Soc.*, 1939, 35, 187-191.  
The authors' experiments were designed to determine the corrosion of metals and alloys exposed to dry sulphuryl chloride (liquid and vapour) and to sulphuryl chloride containing 1 and 4% moisture by volume. In dry sulphuryl chloride, aluminium and nickel were completely corroded. Attack on 15% chromium steel was negligible, while the following were slightly corroded: lead, plate, cast iron, lead, and zinc, the last being appreciably corroded by the vapour. Copper and brass were appreciably corroded, while aluminium bronze and nickel silver showed only traces of corrosion. The presence of moisture in sulphuryl chloride appeared to increase the corrosion of iron, cast iron, bronze, steel, lead, and zinc, and to cause corrosion of aluminium, while corrosion of copper was unaffected.

1742

*Corrosion and Related Phenomena*

MA

The Action of Sulphuryl Chloride on Metals. V.K. Pershko and Kh. L. Tsuytlin (Zhur. Priklad. Khimii (J. Applied Chem.), 1932, 15, (2), 132-137).- (In Russian.) Loss-in-weight determinations were made on a number of metals and alloys exposed to dry sulphuryl chloride (liquid and vapour) and to sulphuryl chloride containing 4.75% moisture by volume. In the dry compound magnesium, aluminium, and nickel were completely resistant. Attack on 1% chromium-steel was negligible, while the following were slightly corroded: boiler plate, cast iron, lead, and zinc, the last being appreciably corroded by the vapour. Copper and brass were appreciably corroded, while aluminium bronze and nickel silver showed only traces of corrosion. The presence of moisture in sulphuryl chloride appreciably increased the corrosion of iron, cast iron, chromium-steel, lead, and zinc, and also caused corrosion of aluminium, while corrosion of copper was unaffected.

--A. S.

1942

MA

4

**Selection of Constructional Materials for Esterification.** V. K. Pershke, E. A. Gerasimov, and S. S. Gerasimov (*Trudy Moskov. Inst. Khim. Mashinostroyeniya* [Moscow Inst. Chem. Eng.], 1939, 17, 104-108; *Khim. Referat. Zhur.*, 1939, 14, 101; *Chem. Abstr.*, 1942, 36, 5451b).—In Russian. Constructional materials resistant to esterification conditions at 120–145°C. and at 125–145°C. were investigated in 3 different media. The compositions of the initial reaction mixture, equal volumes of  $H_2SO_4$  and  $CH_3COOH$ , and at 125–145°C. were investigated in 3 different media. The compositions of these media were approx. equal to those of the media used at various stages of the esterification process: initial mixture, 50%  $CH_3COOH$  in  $H_2SO_4$  solution; Chromium cast iron, EZh 17, EY 4-1, EY 4-2, and chromium-nickel alloys (containing chromium 23%, nickel 14-15%, and molybdenum 3.5-4.5%), copper, tin-lead bronzes (containing 14-16% tin), silicon bronzes, nickel-bronze (containing 30% nickel), and other alloys (corroded in aq.  $H_2SO_4$  solutions to not less than 1-2 cm. annually). Copper alloys corroded considerably more in 50%  $H_2SO_4$ . The corrosion increased sharply in 55%  $H_2SO_4$ . Materials stable in all 3 media were: Thermosolid, Antichlor, and the alloy containing nickel 5% and molybdenum 2%. It is pointed out that steel coils cannot be used in the apparatus. It is recommended to use steel coils preliminarily tinned and uniformly lead-plated with a thick layer of lead, or to change the technical process by using a mixture with a minimum permissible concentration of  $H_2SO_4$ .

1943

17A

\*Investigations on the Protective Sulphate Layer on Lead. V. K. Perekh, Ye. I. Vasyutovich, and Ye. N. Braverman (*Zhur. Priklad. Khim.*, 1939, 11, 1450-1464; *Chem. Zentr.*, 1940, 111, (1), 351).—[In Russian.] The protective film formed on lead in  $H_2SO_4$  can be removed by conc.  $NH_4OH$ ; the use of ammonium acetate and tartrate solutions is not advisable on account of their corrosive action on metallic lead itself. The formation of the film is quickest in 40-78%  $H_2SO_4$ ; in 95-6%  $H_2SO_4$ , no film forms. The protective action of the layer is limited by its porosity. The formation of the layer can be accelerated and its protective action increased by painting the lead surface with  $PbSO_4$ . Addition of antimony, bismuth, tin, copper, zinc, or silver has hardly any effect on the formation of the protective layer. The thickness of the film varies between one and some thousands of molecular layers. The film is easily destroyed in  $H_2SO_4$ , water, and air, and by mechanical action. The layer persists when heated to 85° C., but it is destroyed at 95°. Protection on account of a decrease in adhesiveness and an increase in the solubility of  $PbSO_4$ . A durable film is formed in 67 and 78% nitrosulphuric acid in the presence of 2%  $HNO_3$ , but this is not the case in 95-6% nitrosulphuric acid. Rolled lead sheet is equally or more corrosion-resistant than cut sheet. Addition of 0.05-0.1% cerium to the lead increases its chemical resistance against the action of  $H_2SO_4$ , even at higher temperatures (e.g. 100-115° C.) and high acid concentrations (e.g. 95-6%).



Co

9

The corrosive action of potassium (and other alkali) salt solutions on metals. I. Action on cast iron and lead. V. K. PERSHKEV AND G. I. CHUPAROV. *Ann Inst Polytech Oural* 7, 81-90 (1929-30); *Chem Zvesti* 1931, 1, 2390, cf. C. A. 26, 4433.

The actions of solns. of carnallite, sylvinite, NaCl, KCl and MgCl<sub>2</sub> on cast iron and lead were studied. Cast iron was most strongly attacked by NaCl and least by MgCl<sub>2</sub>, KCl having an intermediate action. Sylvinite soln. corrodes more strongly than carnallite. Dil. solns. are more corrosive than concd. The corrosive action of the former is 1.25-1.75 times that of water, while that of the latter is less than that of water. Satg. the salt solns. with air increases their corrosive action 3 times, light 1.7 times. The lower the content of impurities in the iron, the less it is corroded. A coating of wood tar serves as an effective protection for the iron and lowers the corrosion to 1/10. Lead is 9 times more resistant to the action of salt solns. than iron. M. G. M.

ASB SLA METALLURGICAL LITERATURE CLASSIFICATION

62

| 1ST AND 2ND COPIES                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |
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| <p>4</p> <p>*Constructional Materials for Pans for Drying Organic Dyes. V. K. Dushko and M. A. Voroshilova (<i>Promishlen. Organich. Khimii</i> (Organic Chem. Ind.), 1957, 4, 518-522; C. Abs., 1958, 22, 4788). --[In Russian.] Because of the corrosive action of most of the dyes tested, the use of sheet iron pans for drying is considered impracticable. Excellent results were obtained with aluminium and stainless steel drying pans; copper and bronze are also suitable. The selection of the materials depends on the nature of each dye and on the contaminating products. --S. G.</p> <p>4</p> |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |
| ASD-51A METALLURGICAL LITERATURE CLASSIFICATION                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |        |  |  |  |  |  |  |  |  |  |  |  |  |         |  |  |  |  |  |  |  |  |  |  |  |  |
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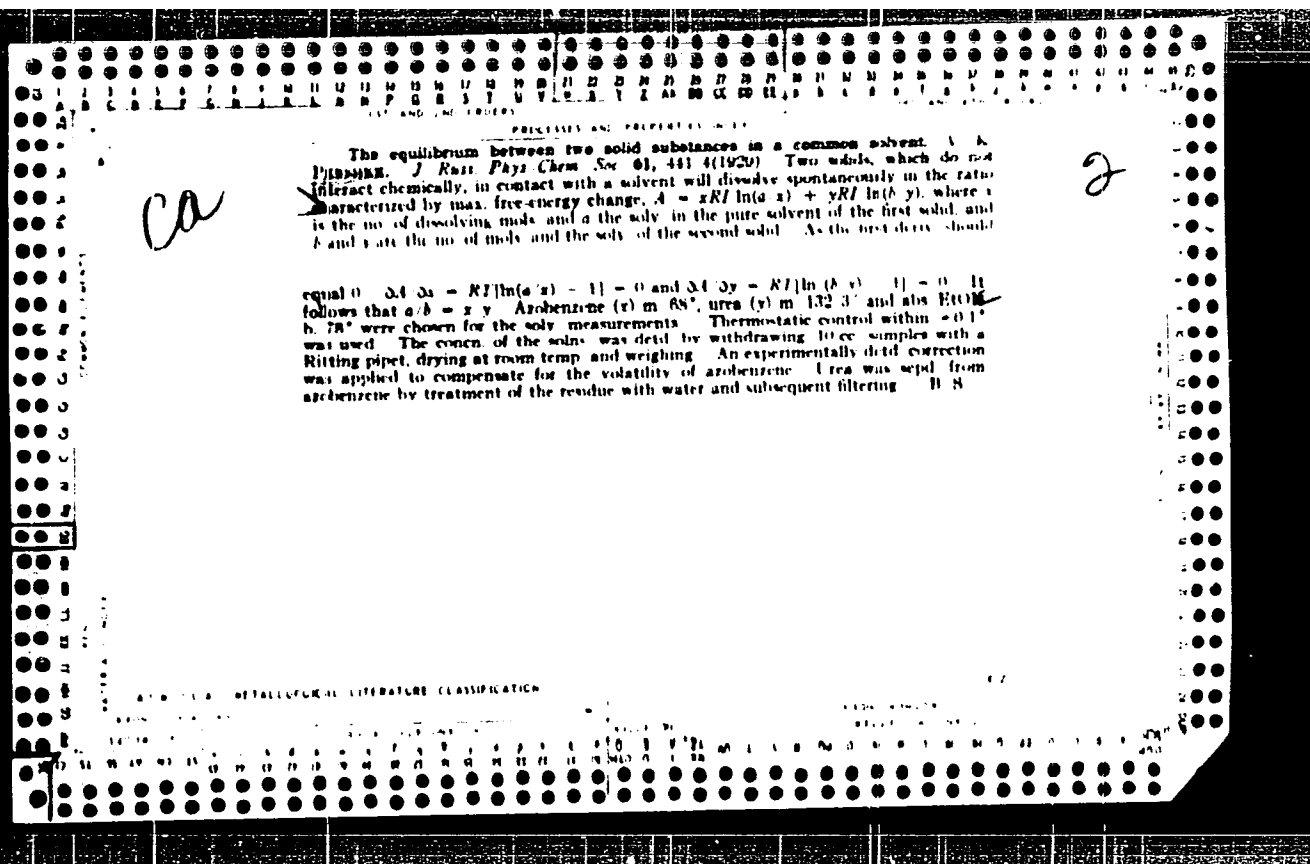
**\*Influence of Metallic Impurities on the Corrosion of Lead in Sulphuric Acid Production.** V. K. Zrinskii (Akimovsk, 1934, 6, 284-287; C.A.B., 1034, 89, 1933).—[In Russian.] Different samples of pure lead (Kahlbaum) alloyed in various proportions with antimony, bismuth, tin, copper, zinc, and silver were exposed to the action of commercial lead-chamber acid containing 1-72% N oxides (based on nitric acid) at 20° C. for 40 hrs. and at 200° C. for 6 hrs. The tabulated results show that bismuth has the greatest corrosive influence, tin somewhat less, antimony and copper very little, if at all, and zinc and silver in the cold have no effect and at 200° C. retard the corrosion. 27 references are given. R. G.

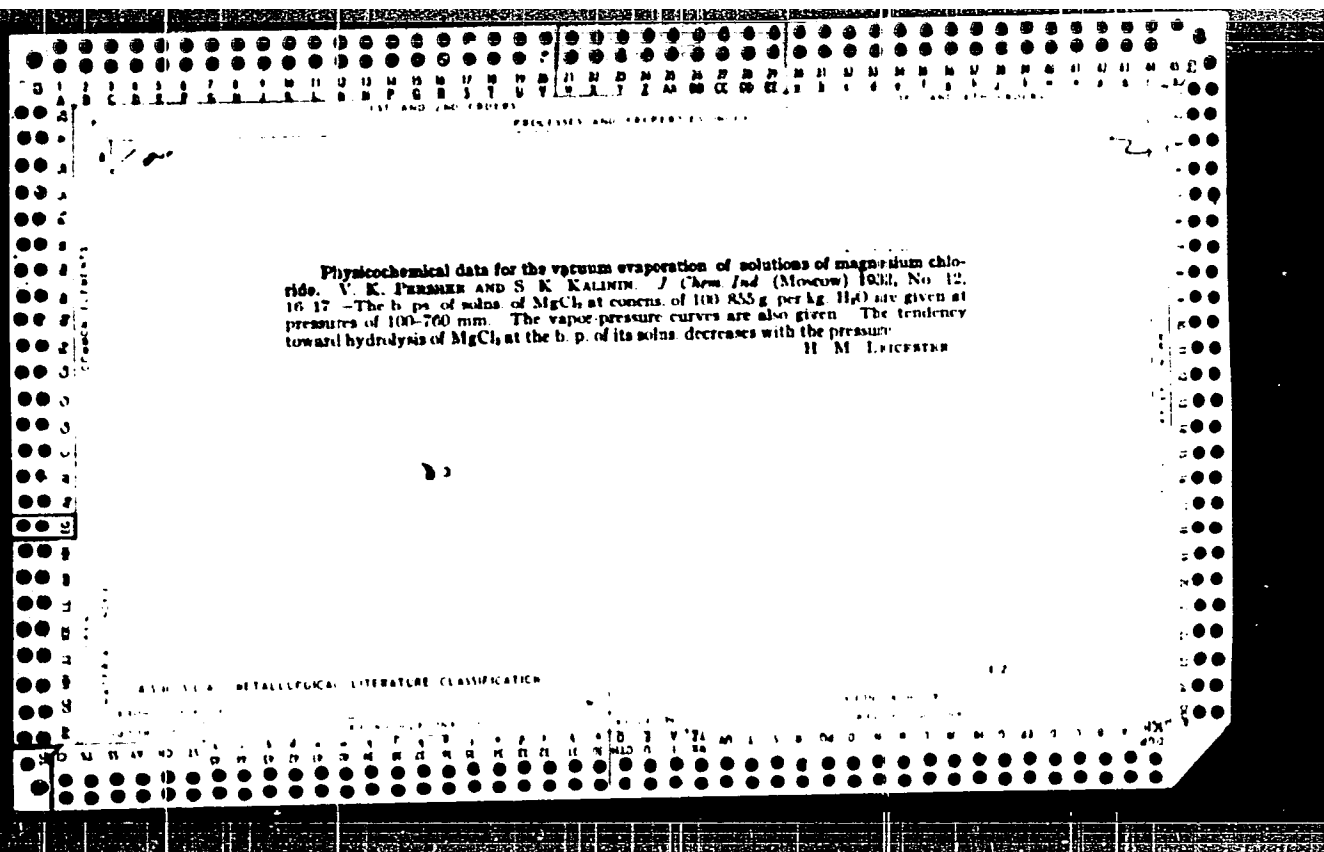
| 1ST AND 2ND COPIES                                                                                                                                                                                                                                                                                                                                                      |  |  |  |  |  |  |  |  |  | 3RD AND 4TH COPIES |  |  |  |  |  |  |  |  |  |
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| PROCESSING AND PROPERTIES INDEX                                                                                                                                                                                                                                                                                                                                         |  |  |  |  |  |  |  |  |  |                    |  |  |  |  |  |  |  |  |  |
| <p>*Corrosion of Metals by Hydrochloric Acid, Chlorine, and Carnallite. V. K. Perel'man and L. A. Peberkina (Khimstud, 1934, 6, 140-141, C. 140, 1934, 66, 5020).- [In Russian.] Iron and copper alloys subjected to the action of gaseous hydrochloric acid and chlorine and carnallite at temperatures of 300°-500° C. showed highly destructive corrosion.—B. G.</p> |  |  |  |  |  |  |  |  |  |                    |  |  |  |  |  |  |  |  |  |
| <p>ASM-AIA METALLURGICAL LITERATURE CLASSIFICATION</p>                                                                                                                                                                                                                                                                                                                  |  |  |  |  |  |  |  |  |  |                    |  |  |  |  |  |  |  |  |  |
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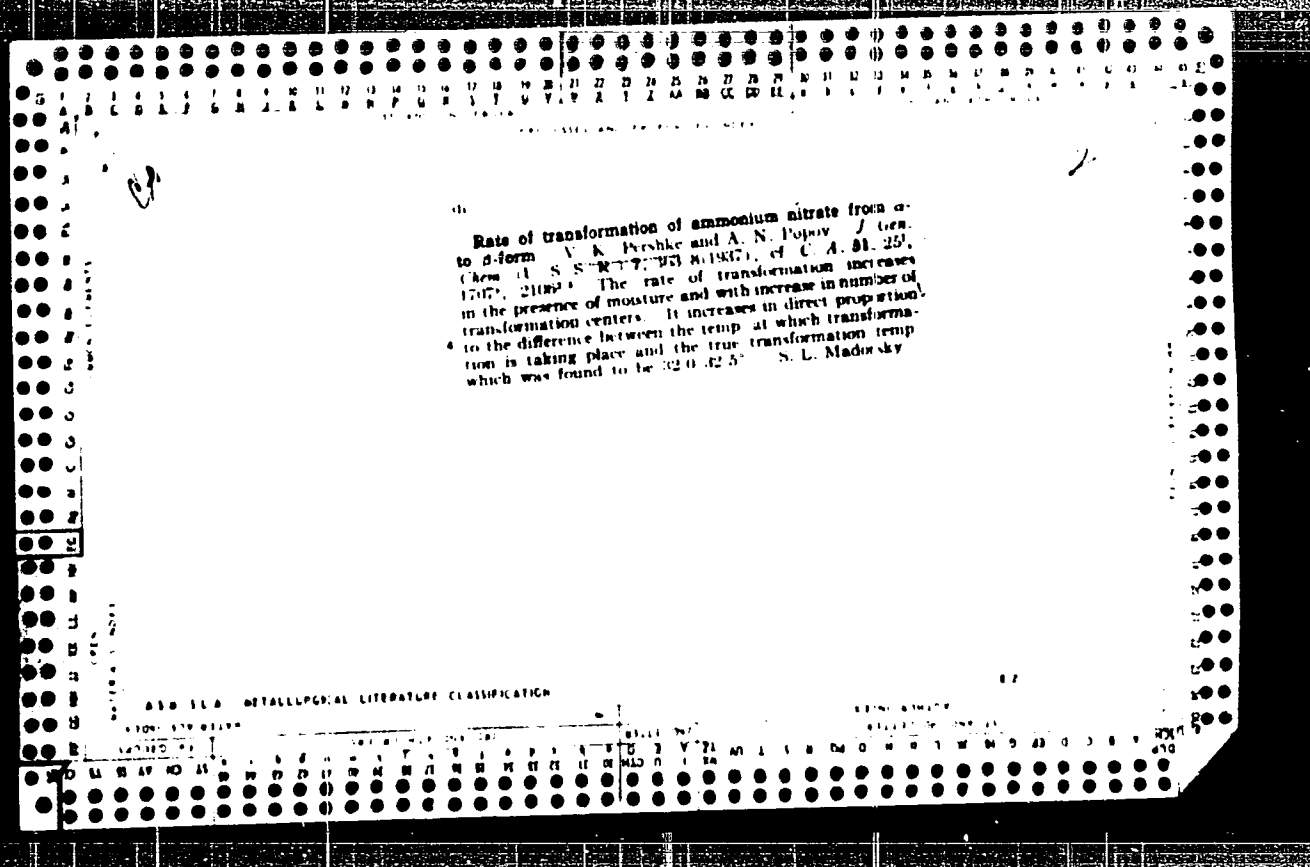
*Influence of Organic Compounds on the Corrosion of Metals by Inorganic Agents.* V. K. Tsvetkov and A. S. Vinogradova (*Promyshlennost' Organicheskoi Khimii* [Org. Chem. Ind.], 1936, 2, 410-423; C. Ab., 1937, 31, 1750).—[In Russian.] A physico-chemical interpretation of the inhibiting action of stable organic compounds and accelerating action of easily reducible organic compounds on the destruction of metals by corroding agents in the process of PhNH<sub>2</sub> production, based on the literature and some experimental data (cf. Jenckel and Braucker, Z. anorg. Chem., 1935, 221, 249; Jimeno and Grifol, Met. Abs., 1936, 3, 276).—S. G.

The heat effect of distribution. V. K. Pershke and A. V. Ishedomonah. *J. Gen. Chem.* (U.S.S.R.) 9, 110-13(1938).—The capil. part of the investigation consisted in the detn. of the heat effect and of the distribution coeff. for the system water-Am alc. (solvent) and ArOH (solute). The substances used were: ordinary distd. water; Am alc. (the 12%-18% fraction); one fraction for each capil.; and 100% special ArOH. Into a 0.5-l. Berzelius flask (in a glass jar and isolated from it by a brass's bath) were introduced 100 ml. of water and layer of cotton) were introduced until a finely 100 ml. of Am alc., and the soln. was stirred with a finely dispersed system was obtained. The usual calorimetric exp. was performed at  $17 \pm 1^\circ$ . The temp. was noted on a Beckmann thermometer every 30 sec. for 10 min. After that 20 g. of ArOH was added and the temp. change detd. for 10 min. The heat capacity of water was taken as 1.000, of Am alc. as 0.554, and of ArOH as 0.484. The mean capil. error was calcd. from  $E = -\sqrt{S/n(n-1)}$ ,  $n = 2$  cal. where  $S =$  sum of the squares of the errors of the individual results,  $n =$  no. of the results. Thus the distribution heat effect at  $17 \pm 1^\circ$  was found to be equal to  $216 \pm 2$  cal. per mol. For the distribution coeff. detn. 100 ml. each of mutually satd. water and Am alc. were placed in a thermostat. To the soln. was added 20 ml. of ArOH and the liquid was shaken for 30 min. After shaking the vol. of both layers were measured; samples were taken from the water layer, and titrated with 0.1 N

NaOH soln. The mean  $K = A/W'$  values obtained at 12, 17, and  $30^\circ$  were 0.913 = 0.003, 0.917 = 0.003, and 0.930 = 0.002, resp. The change of vol.  $\Delta V$  of the solvents during the distribution of ArOH attained 7.5%. The main reason for the change in heat effect with temp. is the change in the mutual sol. in the presence of a third substance. From the capil. results the formula for the heat effect of the distribution of ArOH between water and amyl alc. was found to be  $U = 591.5 - 0.3T + 0.0003T^2$ , and from this a distribution coeff. curve was constructed.  
W. R. Henn







10

Chemical resistance of aluminum in the production of organic compounds. V. K. Darsake and K. B. L. Teltin. *Org. Chem. Ind.* (U. S. S. R.) 6, 682-6 (1939); cf. C. A. 31, 175P.—Several tables are given to show the relative resistance to corrosion of Al in the production of organic compds. and dyes, involving the use of  $\text{HCO}_2\text{H}$ ,  $\text{AcOH}$ ,  $\text{H}_2\text{SO}_4$ ,  $\text{SO}_2\text{Cl}_2$  and other reagents Chas. Blanc



**CA**

The action of suluryl chloride on metals V. K. Ples  
like and Kb. L. Tschlin. J. Applied Chem. (T.S.N.R.)  
12, 182-7 (in French, 1939).—Dry SO<sub>2</sub>Cl<sub>2</sub> does not  
affect Al, Mg and Ni at room temp., but considerably  
corrodes Cu and brass. It corrodes very slightly Cr,  
steel, Fe, cast Fe, Pb, Zn, Al bronze and German silver.  
The corrosion of metals increases in moist SO<sub>2</sub>Cl<sub>2</sub>, especially  
in the case of Zn, Al, cast Fe, Cr steel and Fe. Corrosion  
of Cu and Pb increases comparatively slightly because  
the protective action of the film formed on the metal.

A. A. Podgorny

18

FROM BOMBING  
RELIANT ONE ONE 191

ADD SLA METALLURGICAL LITERATURE CLASSIFICATION

SECTION NO. 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

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THE PREPARATION OF AMMONIUM CHLORIDE FROM THE GASES. V. K. PERSHKE AND K. G. POTASKURV. *J. Chem. Ind. (Moscow)* 1932, No. 11, 15-17.—If gaseous  $\text{NH}_3$  is well mixed with  $\text{HCl}$  from the hydrolysis of  $\text{Mg}$  cement (C. A. 25, 4077) and the resulting  $\text{NH}_4\text{Cl}$  drawn through a chamber at  $130-70^\circ$  at a rate of 20 cc. per min., 75% settles in this chamber. The finely dispersed particles are then drawn into a chamber at room temp., where practically all the remainder is caught as a satd. soln. by the condensed  $\text{H}_2\text{O}$  of the process.

H. M. LEICHTER

ASD S.L.A. METALLURGICAL LITERATURE CLASSIFICATION

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REMARKS

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| 1ST AND 3RD COLUMNS                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |  |  |  |  |  |  |  |  |  | 2ND AND 4TH COLUMNS |  |  |  |  |  |  |  |  |  |
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| PROCEDURES AND PROPERTIES INDEX                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 |  |  |  |  |  |  |  |  |  |                     |  |  |  |  |  |  |  |  |  |
| 6C                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |  |  |  |  |  |  |  |  |  | 8 I 8               |  |  |  |  |  |  |  |  |  |
| <p><b>Hydrochloric acid and magnesia from magnesia chloride.</b> V. K. PASHKIN and G. I. CHUFANOV (J. Chem. Ind., Russia, 1930, 7, 332-335).—The solid prepared from 35% <math>MgCl_2</math> solution (100 pts.) and <math>MgO</math> (10 pts.) is dried at <math>100^\circ</math> to 80% loss in wt., and calcined at <math>500^\circ</math> in a mixture of air and <math>H_2O</math> vapour (1 pt. <math>H_2O</math> to 1 pt. total <math>MgO</math>). The <math>MgO</math> contains 0.5% Cl.</p> <p>CHEMICAL ABSTRACTS.</p> |  |  |  |  |  |  |  |  |  |                     |  |  |  |  |  |  |  |  |  |
| <p>ASAC-SEA DETALLURGICAL LITERATURE CLASSIFICATION</p> <p>33000 31000000</p> <p>330000 31000000</p> <p>330000 31000000</p>                                                                                                                                                                                                                                                                                                                                                                                                     |  |  |  |  |  |  |  |  |  |                     |  |  |  |  |  |  |  |  |  |



**Destructive action of phosphoric acid on metallic alloys.** V. K. LAMHAR, M. P. KAPISHIN and E. V. YUSHMANOV. *J. Chem. Ind. (Moscow)*, 9, 855-5 (1931). Expts are described on the resistance of cast Fe and Cu alloys contg. Sn, Zn, Al, Pb and Ni to the action of 84.7%  $H_3PO_4$  at a temp. from  $20^\circ$  to  $150^\circ$ . The results are given in tables. Alloys contg. 78% Cu are resistant to  $H_3PO_4$ , with 83-80% Cu they are satisfactory but with a Cu content lower than 83% they are not resistant to  $H_3PO_4$ . Alloys with the same Cu content and nearly equal percentages of Pb, Al and Sb show the same resistance. With the same Cu content, Ni instead of Zn and Pb, or Sn and Zn and Pb instead of Zn increase the resistance; Pb instead of Sn has no influence. Pb and Sb instead of Zn do not change the resistance of the alloy. Alloys with a high content of Pb and Zn are not resistant. Brass with a Cu content of 83-90% and Cu-Ni alloys with 80% Cu are recommended for app. coming in contact with concd.  $H_3PO_4$  at  $150^\circ$ . (C)

CP

6

Preparation of dehydrated magnesium chloride from magnesium cement. *Verkhovskiy and A. P. Lohvitskaya. Kzhu (U. S. S. R.) 1933, No. 4, 33-5. — A method for prepn of dehydrated  $MgCl_2$ .  $Mg$  cement in briquets (prepd from a 35% soln of  $MgCl_2$  and  $MgO$  (10 g. per 100 g. soln)) were dried after solidifying in an elec. drier at 150°. Then the temp. was raised to 400-600° and the gas mixt. (air- $HCl$ ) was drawn through the briquets by suction. A product contg. about 97%  $MgCl_2$  was obtained.* James Borrel

AND SEE DETAIL OF LITERATURE CLASSIFICATION

**The use of nickel chlorinator.** A. K. Pershke and M. L. Gligorov, *Chem. Abstr.* 1954, 48, 8785 (1954).

The corrosion rates of com. Ni, Ag, Pb and Cu, Si and Cr-cast iron and steel by the gas and liquid naves in the direct chlorination of org. compounds were studied by suspending weighed, polished plates (15 sq. cm.) from a rubber sleeve and inside the liquid reaction vessel. The corrosion rate was calculated from the loss in weight in mg./hr. In the chlorination of 100% AcOH with the Si catalyst at 100-200 for 1-4 hrs. only Ag can be successfully used. Cr-steel and cast iron inhibit the formation of  $\text{CH}_3\text{COCl}$ . In the presence of 2-4 MeC<sub>2</sub>H<sub>4</sub>ClSO<sub>4</sub>H from p-MeC<sub>2</sub>H<sub>4</sub>SO<sub>4</sub>H in 100% H<sub>2</sub>SO<sub>4</sub> at 10-5 for 10 hr. Ni is nearly 100% resistant to the action of gases and is somewhat more stable to the action of H<sub>2</sub>SO<sub>4</sub> than cast iron, the corrosion of which progressively increases, and a little less stable than Cr-cast iron and steel. In the chlorination of dry p-ClC<sub>2</sub>H<sub>4</sub>Me at 100-200 for 30-40 hrs. Ni proved to be the most stable metal. Ni, Ag and glass app. give chiefly ClC<sub>2</sub>H<sub>4</sub>Cl, while cast iron, Cu and Pb give considerable chlorination product substituted in the nucleus. In the presence of water Ni cannot be used for chlorination and bromination. Its corrosion was by gases 16.2 and by liquid 17.2 g./sq. m./hr. after 9 hrs. of interaction. Satisfactory results in the presence of water were obtained with Si-cast iron. Chris. Blanc.

Chemical resistance of chromium and aluminum steels to sulfur vapors at high temperatures V. K. Pershke, S. S. Gostev and I. V. Oknin. *Khim. Mashinostroeniye* 6, No. 4, 27-30 (1967); *Chem. & Industry* 40, 265-6. Up to 700°C, "fechtal" is a ferritic alloy contg. 12-14% Cr, 3-5% Al and 0-15% C and especially chromal exhibit satisfactory resistance to S vapors (0.3 g./per l.), whereas materials such as boiler plate, Cr steel, Ni steel, etc., under the same conditions undergo such strong corrosion as to render them useless. S corrosion is of considerable importance in the corrosion of "fechtal." On the other hand, variation in the content of H<sub>2</sub>S up to 0.5%, of SO<sub>2</sub> up to 4%, and of CS<sub>2</sub> up to 7% has no effect. Chromal is twice as resistant as "fechtal." The optimum proportions of the alloying metals are Cr 13-14 and Al 4-5%. A. P. C.



| 1ST AND 2ND COLUMNS                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 3RD AND 4TH COLUMNS |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
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| PROCESSES AND PROPERTIES INDEX                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |                     |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| <p><i>A</i></p> <p>Influence of metallic impurities on the corrosion of lead in sulfuric acid production. V. K. Pershke. <i>Khimicheskii Zhurnal</i>, 284 7(1034).—Different samples of pure Pb (Kahlbaum) alloyed in various proportions with Sb, Bi, Sn, Cu, Zn and Ag were exposed to the action of factory lead-chamber acid contg. 1.72% N oxides (based on HNO<sub>3</sub>) at 20° for 40 hrs. and at 240° for 0 hrs. The tabulated results show that Bi has the greatest corrosive influence, Sn somewhat less, Sb and Cu very little, if at all, and Zn and Ag in the cold have no effect and at 240° retard the corrosion. Twenty-seven references. Chas. Blum.</p> |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |                     |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| <p>WITH S.E.A. METALLURGICAL LITERATURE CLASSIFICATION</p> <p>1970-1971</p>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |                     |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

CO

Preparation of dehydrated magnesium chloride by means of ammoniac. V. K. Pershke and S. K. Kalinin. Khol 1933, No. 6, 17-21. —Lab. expts. were made to verify Brit. pat. 330,650 (C. A. 26, 5048) for the prepa. of dehydrated  $MgCl_2$ .  $NH_3$  gas was passed through a soln. of  $MgCl_2$  contg.  $NH_4Cl$  to form the ammoniate ( $MgCl_2 \cdot 6NH_3$ ,  $MgCl_2 \cdot 2NH_3$ ;  $MgCl_2 \cdot NH_3$ ). The best results were obtained with a soln. of 17.5%  $MgCl_2$  and 6%  $NH_4Cl$  at 20° when practically all of the  $MgCl_2 \cdot 6NH_3$  is pptd out.  $MgCl_2$  of 95-96% dehydration was obtained. James Sorrel

ASD 35.4 DETAILING LITERATURE CLASSIFICATION

PERSHKE, V. K., LAKEDMONSKIY, A. V.

"On the Thermal Effect of Distribution," Zhur. Obshch. Khim., 9 No. 2, 1939.  
Laboratory of Physical Chemistry, Moscow Institute of Nonferrous Metals and Gold.  
Received 27 May 1938

U-1517, 22 Oct 1951

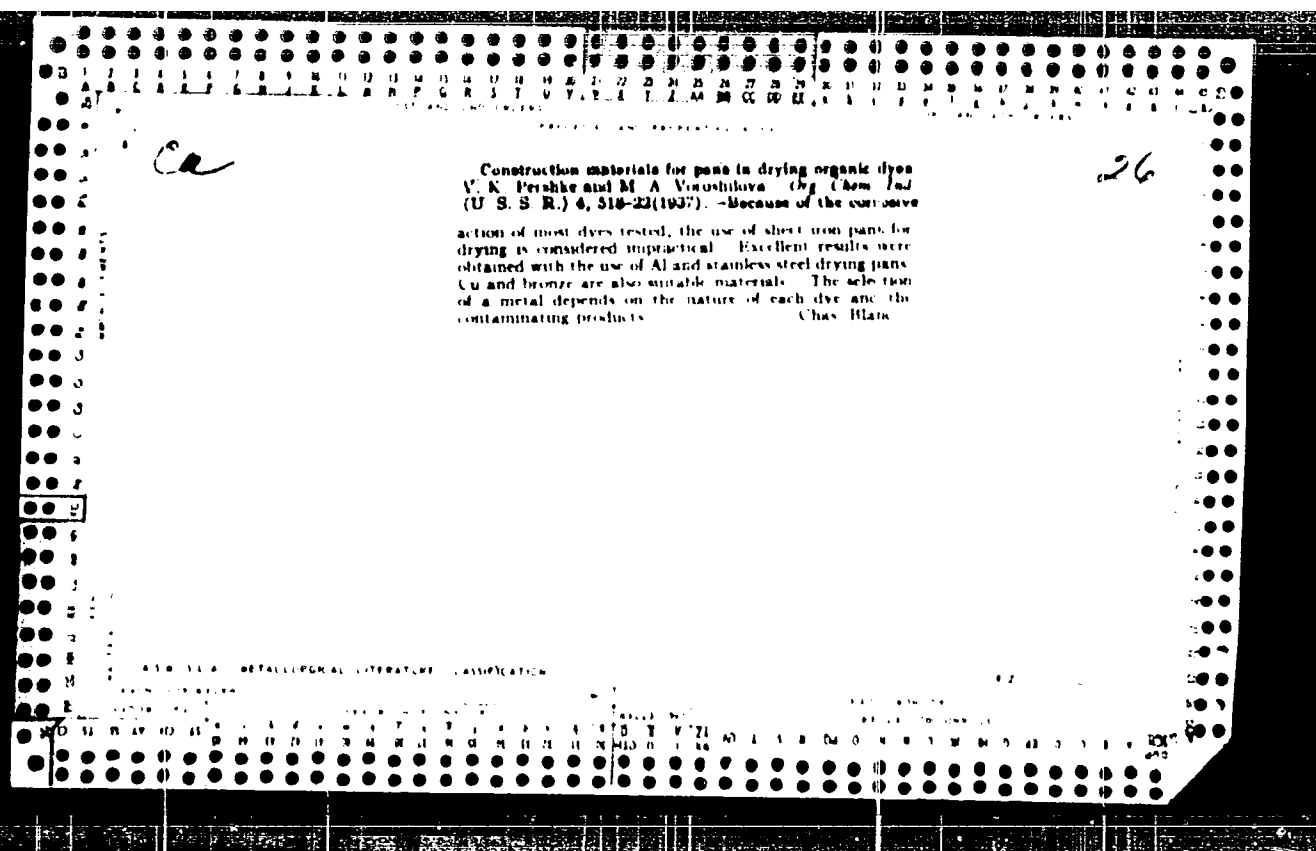
**Corrosion of metals by hydrochloric acid, chlorine and carnallite.** V. K. Pershke and L. A. Pecherkins. *Khim. zvezd* 6, 140 (1974). Fe and Cu alloys subjected to the action of gaseous HCl and Cl<sub>2</sub> and carnallite at temps. of 200-700° showed highly destructive corrosion. © 1975

**APPROVED FOR RELEASE: 06/15/2000**

**CIA-RDP86-00513R001240120010-0"**

*m*

**The Use of the Nickel Chlorinator.** (Corrosion of Nickel, Silver, etc.)  
V. K. Ivashko and M. G. Grigor'ev (*Primen. Organ. Khimii* (*Organic Chem. Ind.*), 1958, 5, (1), 50-53; *C. Abstr.*, 1958, 52, 6218).—[In Russian.] The corrosion rates of commercial nickel, silver, and a number of ferrous materials by the gas and liquid mixtures involved in the chlorination of organic compounds were studied by suspending weighed, polished plates (15 cm.<sup>2</sup>) from a stirrer above and within the liquid reaction mixture. The corrosion rate was calculated from the loss in weight in gm./m.<sup>2</sup>/hr. In the chlorination of 100% AcOH with the  $SnCl_4$  catalyst at 105°-150° C. for 15-17 hrs., only silver can be successfully used. In the preparation of 2:4-dichloro-1,3,5-trimethylbenzene from p-MeC<sub>6</sub>H<sub>4</sub>SO<sub>3</sub>H in 100% H<sub>2</sub>SO<sub>4</sub> at 40°-45° C. for 49 hrs., nickel is almost 100% resistant to the gases and is somewhat more stable to the action of H<sub>2</sub>SO<sub>4</sub> than is cast iron, the corrosion of which progressively increases, and a little less stable than chromium cast iron and steel. In the chlorination of dry p-ClC<sub>6</sub>H<sub>4</sub>Me at 155°-220° C. for 30-40 hrs., nickel proved to be the most stable metal. In the presence of water nickel cannot be used for chlorination and bromination. Its corrosion was by gases 16-2 and by liquid 17-2 gm./m.<sup>2</sup>/hr. after 9 hrs. of interaction. Satisfactory results in the presence of water were obtained with silicon cast iron.



**Influence of aromatic acids on the corrosion of metals by sulfuric acid** V. K. Pecher and R. I. Grayskaya (*Ing. Chem. Ind.* (U.S.S.R.) 5, 58 (1968), (U.S.S.R.) 31, 176P). The inhibiting action of aromatic acids on the corrosion of boiler plate (0.01% C), Cr steel (10.0% Cr and 0.24% C) and com. Pb by 10, 40 and 70%  $H_2SO_4$  at 20, 60 and 90° was studied by stirring polished metal plates of equal wt. and dimensions for 3-6 hrs. in  $H_2SO_4$ , with and without the addition of 1% of org. acids, and in 1% aq. solns. of the org. acids. The tabulated results show that the inhibiting action of salicylic, benzoic and phthalic acids is about equal and considerably lower than that of *p*-toluene-*o*-benzoic acid. In the presence of the org. acids the corrosion of Fe and steel is retarded at max. 40%  $H_2SO_4$  and 90° (cf. Jenckel and Braucker, *C. A.* 20, 2428), very little, if at all, at higher  $H_2SO_4$  concns. and higher temps., while at the concn. of 70%  $H_2SO_4$  the rate of corrosion is slightly accelerated. In general, the rate of corrosion is influenced more by the  $H_2SO_4$  concn. than the temp. Though the addition of org. acids reduces the corrosion considerably, the abs. value of the corrosion by  $H_2SO_4$  is so great that the inhibiting effect of the addition of org. acids has no practical value. The corrosion of Pb by org. acids and  $H_2SO_4$  is comparatively small at 20° and 60° and considerable at 90°. Chas. Blanc

ASB-I.L.A. METALLURGICAL LITERATURE CLASSIFICATION

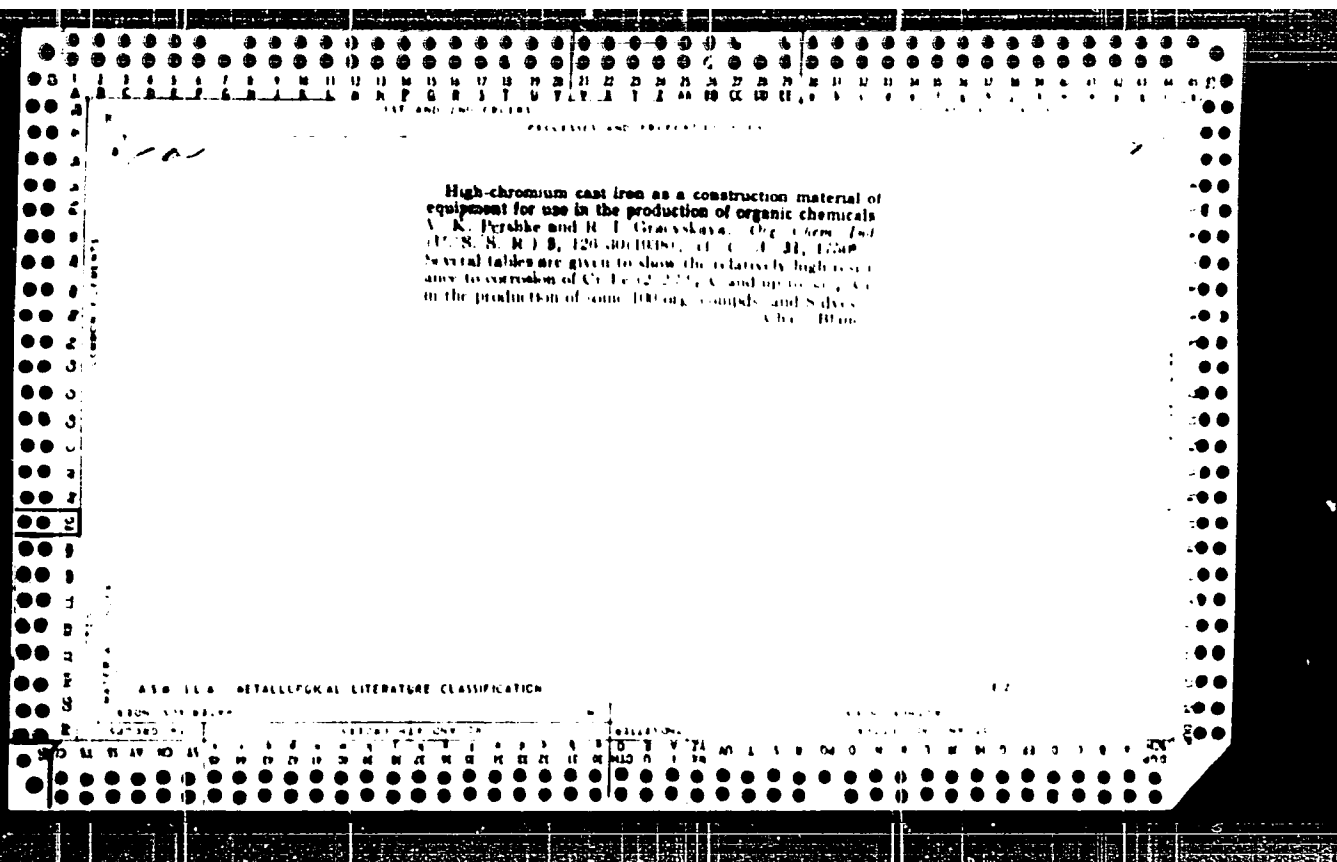
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101

PROCESSING AND PREPARATION

Influence of organic matter on the corrosion of metals by mineral agents. V. K. Pershke and M. A. Voroshilova. *Org. Chem. Ind. (U.S.S.R.)* 5, 562 (1968), cf. C. A. 31, 1754p; 32, 4788p. Addnl. data on the accelerating action of aromatic nitro compds. on the destruction of metals by corroding mineral agents are given. Chas. Blanc

ASD SLA METALLURGICAL LITERATURE CLASSIFICATION

101

[illegible]

**Corrosion of metals in the amination of aromatic chlorides** V. K. Pershke and Kh. L. Fakhim (*Chem. Ind. U. S. S. R.* 5, 1947, 1948, 1; C 4 31, 1750). Cr-Ni, Cr and Cu steels and boiler plate resist the corrosive action of  $\text{NH}_4\text{Cl}$  and  $\text{CuSO}_4$  (catalyst) in the production of  $\beta$ -aminoanthraquinone from the Cl deriv. and concd.  $\text{NH}_4\text{OH}$  at 200° and 50 atm. The inhibition of the corrosion of Fe in the presence of excess  $\text{NH}_4\text{OH}$  and gaseous  $\text{NH}_3$  is traced to the impeded disson of the  $\text{NH}_4\text{Cl}$  and copper-aminoma complex formed in the reaction.

Chas Blanc

(Chas. Blanc)

ASD SLA DETALLURGICAL LITERATURE CLASSIFICATION

**CIA-RDP86-00513R001240120010-0"**

CA

Investigation of the chemical resistance of manganese and antimony bronzes. V. K. Pershke and S. I. Vasyutovsk. *Trudy Akad. Nauk SSSR, Metalloved.* 1939, No. 7, 109-12. *Khim. Referat. Zhur.* 1940, No. 1, 142. The chem. stability of Mn and Sb bronzes for the purpose of replacing Sn bronze was investigated. Common diagrams of Cu-Mn and Cu-Sb alloys are described. Alloys contg. not more than 7% of Mn and 4-5% of Sb were prepd. on the base of these alloys. Bronzes of such compns. are solid solns. Up to 2% of Sn was added to Mn bronze. Various concns. of  $H_2SO_4$  at 70-100° were used as the corrosion medium. Al and Sb bronzes were examd. for comparing their properties with those of Mn and Sn bronzes. The Sb-, Mn- and Mn-Sb bronzes, being solid solns. at the concns. investigated, are stable in concd.  $H_2SO_4$  solns. They are slightly less stable against  $H_2SO_4$  than the Al- and Sn bronzes. Sb bronzes are slightly less stable than other bronzes and their mech. properties are poorer than those of other bronzes. Mn-Sb and Mn bronzes are equally stable. The addn. of Sn is, therefore, not recommended. The temp. limit of the stability of Mn bronzes in 60-70%  $H_2SO_4$  is 80°. W. R. Horn

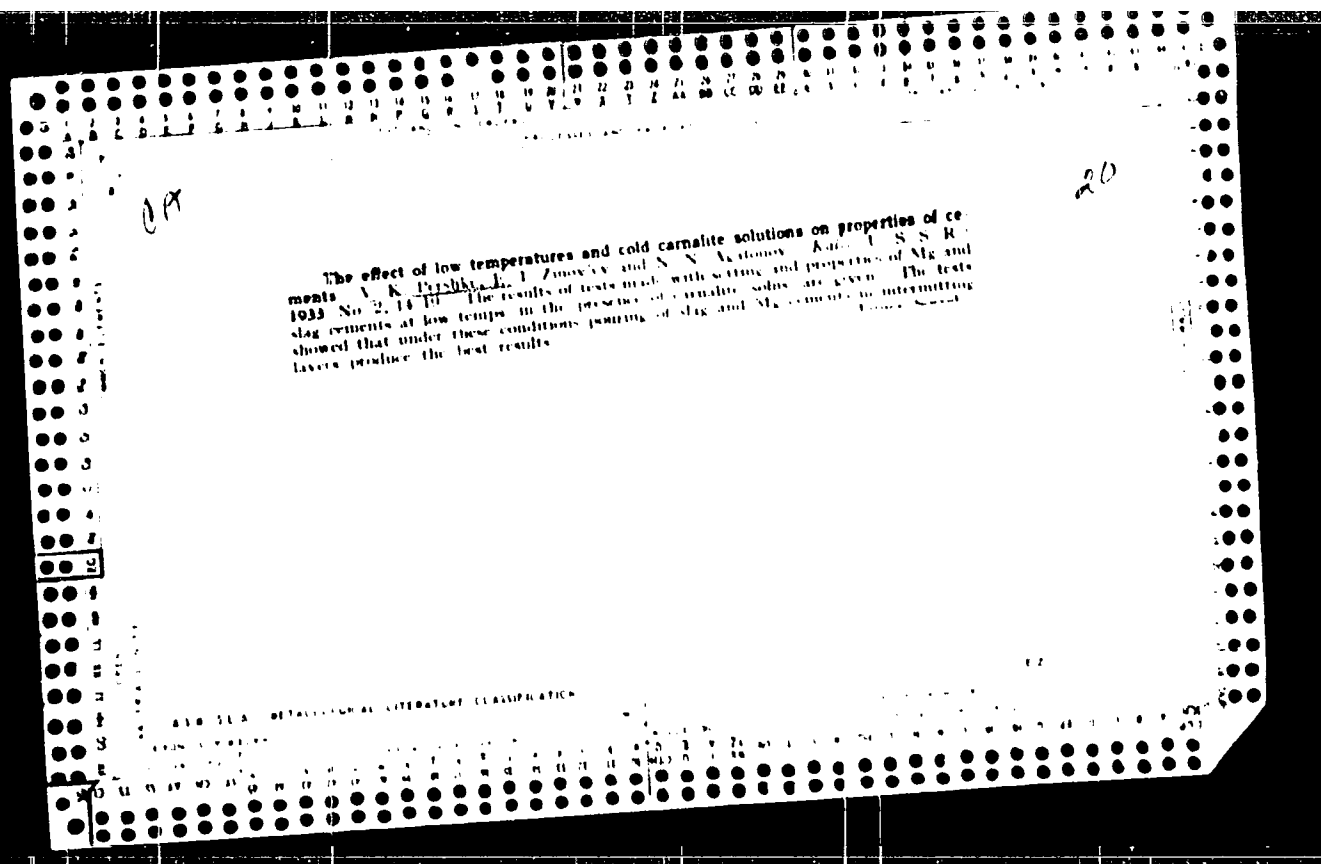
ADD SLA DETAIL LITERATURE CLASSIFICATION

CA

Selection of construction materials for esterification  
V. K. Porabko, I. V. Okun and S. S. Gostev. *Trudy  
Mashin. Inst. Khim. Mashinostroyeniya* 1939, No. 7, 103.  
Khim. Referat. Zhur. 1940, No. 6, 144.— Construction  
materials (20 metals and alloys) stable under esterifica-  
tion conditions at variable compns. of the initial reaction  
mixture (equal vols. of  $H_2SO_4$  (1.8) and 90% alc.) and at 125  
45° were investigated in 3 different media. The compns.  
of these media were approx. equal to those of the reaction  
media at various stages of the esterification process (ini-  
tial mixt., 80%  $H_2SO_4$  and 20%  $H_2SO_4$  soln.). Cr cast  
Fe, RZh-17, EYa-1, EYa-Ti, Fe-Cr-Ni alloys (contg. Cr  
23.5, Ni 14.15 and Mo 3.5 + 5%), Cu, Sn-Pb bronzes  
(contg. from 11 to 30% of Pb), Sn bronzes, Ni bronze  
(contg. Ni 30%) and other alloys corroded in aq.  $H_2SO_4$   
solns. to not less than 1.2 cm. annually. Alloys  
on Cu base corroded considerably more in 80%  $H_2SO_4$ .  
The corrosion decreased sharply in 55%  $H_2SO_4$ .  
Materials stable in all 3 media were: thermosolid, anti-  
chlor and the alloy contg. Ni 50 and Mo 50%. Inasmuch  
as steel coils cannot be used in the app. P., O. and G.  
propose either to use steel coils preliminarily tinned and  
uniformly Pb plated with a thick layer of Pb or to change  
the tech. process by using a mixt. with a min. permissible  
concn. of  $H_2SO_4$ .  
W. R. Helm

AS-8-51A METALLURGICAL LITERATURE CLASSIFICATION

| SECTION | SUBSECTION | CLASSIFICATION | REMARKS |
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| 1ST AND 2ND ORDERS                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          |  |  |  |  |  |  |  |  |  | 3RD AND 4TH ORDERS                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |  |  |  |  |  |  |  |  |  |
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| PROCESSES AND PROPERTIES INDEX                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |  |  |  |  |  |  |  |  |  |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        |  |  |  |  |  |  |  |  |  |
| <p><b>Investigation of the protective sulfate film formed on lead (by the action of sulfuric acid).</b> V. K. Perabke, E. F. Vasyutovich, E. N. Braverman and O. K. Varentsova. <i>J. Applied Chem. (U. S. S. R.)</i> 11, 1450-54 (in French, 1954) (1958). — The protective film formed by dipping lead into <math>H_2SO_4</math> of any concn. up to 95.6% can be removed by treatment with concd. <math>NH_4OH</math> soln. <math>NH_4OAc</math> and <math>NH_4</math> tartrate cannot be used, because they dissolve the lead. The best concn. of <math>H_2SO_4</math> for the formation of film was 60-70%. The film did not completely protect Pb</p> |  |  |  |  |  |  |  |  |  | <p>from corrosion, apparently because of porosity; however, the corrosion under the film was very slight. The formation of film and its protective effect can be accelerated and improved by rubbing <math>PbSO_4</math> into the Pb surface. <math>PbSO_4</math> stopped up pores and crevices in the film and adhered to the surface of the metal. Addn. of other metals (Sb, Bi, Sn, Zn, Fe and Cu) had no effect on film formation and, hence, on corrosion. The thickness of film varied from one to several thousand mols. The film was easily destroyed in water and in air and by various mech. treatments. A temp. of 85° did not affect the formation and existence of film, but at 95° the film was destroyed and renewed formation failed. The film was formed and existed in 67% (better) and in 78% nitrosylsulfuric acid; addn. of 2% <math>HNO_3</math> did not change the corrosive effect. The film did not form or exist in 95.6% nitrosylsulfuric acid. Addn. of 0.03-0.1% Fe considerably improved the chem. stability of Pb against <math>H_2SO_4</math> at high temp. and concn. of acid. Thus, the stability of Pb increased 24-45 times in 78% <math>H_2SO_4</math> at 100-110° and 4 times in 95.6% acid. A. A. Podgorov</p> |  |  |  |  |  |  |  |  |  |
| <p>ASB-514 METALLURGICAL LITERATURE CLASSIFICATION</p>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      |  |  |  |  |  |  |  |  |  |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        |  |  |  |  |  |  |  |  |  |

137 AND THE 800173 PROCESSING AND DOCUMENTING NOTES

bc

B-I-8

Influence of metallic impurities on corrosion of lead by sulphuric acid production. V. K. Ponomarev (Khimistr., 1934, 6, 266-267). Samples of Pb containing impurities were exposed to  $H_2SO_4$  (1.79% of  $N$  oxide) at 20° (80 hr.) and at 300° (8 hr.). Bi had the most corrosive influence, Sn had less, Sb and Cu had little action, and Zn or Ag had no effect in the cold, but retarded corrosion at 300°.

On. Ann. (r)

ASS. SLA METALLURGICAL LITERATURE CLASSIFICATION

FROM SOURCE

137 AND THE 800173



18

Hydrochloric acid and magnesia from magnesium chloride. V. K. PERSHIK AND G. I. CHUTAROV. *J. Chem. Ind. (Moscow)* 7, 332-5 (1950). A method is described for  $MgCl_2$  decomposition to  $MgO$  and  $HCl$  by prep. of solid cement from 45%  $MgCl_2$  soln. and  $MgO$  (two parts  $MgCl_2$  soln. to 10 parts  $MgO$  by wt.), drying at 200° to 50% loss of wt., calcining at 500° in air water vapor mist (1 part vapor to 1 part  $MgO$  by wt. based on total  $Mg$ ). The products obtained are  $HCl$ , sp. gr. 1.14-1.15, and  $MgO$  containing 0.5%  $Cl$ . JAMES CURRIE.

DETAILS OF LITERATURE CLASSIFICATION

Influence of organic compounds on the corrosion of metals by inorganic agents. V. K. Fritshchik and A. S. Vinogradova. (Dg. Chem. Ind. (U. S. S. R.) 2, 410-23 (1959)). A physicochem. interpretation of the inhibiting action of stable org. compds. and accelerating action of easily reducible org. compds. on the destruction of metals by corroding agents in the process of  $\text{PhNH}_2$  production is based on literature and some expl. data (cf. Jiracki and Bräcker, C. A. 30, 2428; Jimeno and Giffol, C. A. 30, 3378, 4133).

ASD SLA METALLURGICAL LITERATURE CLASSIFICATION

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| 1ST AND 2ND COPIES                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |  | 3RD AND 4TH COPIES |  |
| <p><b>BC</b></p> <p><b>Corrosive action of potassium (and other alkali) salt solutions on metals. I. Cast iron and lead. V. K. Ponomarev and G. I. Chudakov (Ann. Inst. Polytech. Omsk, 1958-1960, 7, 81-90).—Cast Fe was most strongly attacked by NaCl, less by KCl, and least by MgCl<sub>2</sub>; sylvite is more corrosive than carnallite, and dil. solutions are more corrosive than conc. solutions. Light (1.7 times) and air (3 times) increase the action. Pure Fe is less corroded than impure; Pb is "9 times more resistant than Fe."</b></p> <p><b>B-I-4</b></p> |  |                    |  |
| <p>450-51.0 METALLURGICAL LITERATURE CLASSIFICATION</p>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         |  |                    |  |
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PERSHKHAVA, A.A.

Case of acute appendicitis in a newborn infant. *Pediatrics* 38  
no.10:77-78 0 '60. (MIRA 13:11)

1. Iz rayonnoy bol'nitsy (glavnyy vrach A.A. Pershkhava) sela  
Orshanka Mariyskoy ASSR.  
(APPENDICITIS) (INFANTS (NEWBORN)---DISEASES)

PERSHKHOVA, A. A.

PERSHKHOVA, A. A.

Early hospitalization of emergency surgical cases in rural areas.  
(MIRA 7:11)  
Khirurgiya no.8:58-60 Ag '54.

1. Iz Orshanskoy rayonnoy bol'nitsy Mariyskoy ASSR (glavnyy vrach  
T.Yu.Pirogova)

(RURAL CONDITIONS.

early hospitalization of emergency surg. cases)

(EMERGENCIES.

early hospitalization of emergency surg. cases in rural  
cond.)

PERSHMAN, R.E.

NEUR

1666. Selective Involvement of the Hand in Parasagittal Meningioma. (Избирательное поражение руки при парасагитальных менигиомах)

R. E. PERSHMAN. Вопросы Нейрохирургии [Vop. Neirokhir.] 14, No. 2, 21-27, March-April, 1950.

Symptoms or signs present in the hand are at times the only, or the main, localizing features in cases of parasagittal meningioma. They were observed in 16 cases out of 70 in this series. Ten such cases are described and the author suggests that these phenomena are produced in different ways. Pressure on the motor pathways in the subcortical substance is one of them, and the irritative action of the tumour on contiguous areas, which are themselves in a state of "altered function", is another. The cause in most cases, however, is interference with venous drainage from the lower part of the parietal lobe through the superficial cortical veins, which are compressed by the tumour at a point near their entry into the sinuses.

L. Crane

See also Section Radiology, Abstract 1435.

Abstracts of World Medicine  
Vol 8 1950

PERSHMAN, R.E.K.

25307 PERSHMAN, R.E.K. Voprosy O Mikhloys-Epilepsii. Nevrologiya i  
Psiknatriya. 1949, No. 4, S. 30-42

SO: Letopis' No. 13, 1949

PERSHMAN, R. Ye., kand.med.nauk

Diagnostic errors in craniopharyngiomas. Probl.sovr.neirokhir.  
3:259-267 '59. (MIRA 16:6)  
(PITUITARY BODY—TUMORS)



PERSHMAN, R. Ye.

Cand. Med. Sci.

Dissertation: "Neurologic-Clinical Treatment of the Primary Extracerebral  
Tumors of the Parasagittal Region."

27/6/50

Acad. Med. Sci. USSR

SO Vecheryaya Moskva  
Sum 71

PERSHMAN, R.Ye.; KONOVALOV, Yu.V.

~~\_\_\_\_\_~~  
Clinical aspects of so-called "recurrent" astrocytomas of  
the cerebellum in children. Vop.neirokhir. 19 no.5:20-25  
S-O '55. (MLRA 8:11)

1. Iz Nauchno-issledovatel'skogo ordena Trudovogo Krasnogo  
Znameni Instituta neyrokhirurgii imeni akad. N.N.Burdenko  
Akademii meditsinskikh nauk SSSR.

(ASTROCYTOMA,  
cerebellum, recur. in child.)

(CEREBELLUM, neoplasms,  
astrocytoma, recur. in child.)

BURGMAN, G.P.; VOZNAYA, A.TS, NITROVANOVA, N.P.; PERSHMAN, R.Ye.

Preoperative and postoperative cerebrospinal fluid in cerebellar medulloblastomas and its clinical significance. Vop.neirokhir. 19 no.6:25-32 N-D '55. (MIRA 9:1)

1. Iz nauchno-issledovatel'skogo ordena Trudovogo Krasnogo Znameni instituta neyrokhirurgii imeni Akad. N.N.Burdenko Akademii Meditsinskikh nauk SSSR.

(CEREBROSPINAL FLUID, in various diseases, medulloblastoma of cerebellum)

(MEDULLOBLASTOMA, cerebellum, CSF in)

(CEREBELLUM, neoplasms, medulloblastoma, CSF in)

